CLAIMS

An anthraguinone compound of the general formula I or a salt 1. thereof

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$$R^4$$
 R^3
 R^2
 R^2

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in which R1 to R4 are each selected from the group consisting of H, C1-4 alkyl, X^1 , -NHR⁰N (R⁵)₂ in which R⁰ is a C₁₋₁₂ alkanediyl and each R⁵ is H or optionally substituted C₁₋₄ alkyl, and a group of formula II

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$$-NH-R^{0}-N \xrightarrow{R^{0}} R^{6}$$

$$R^{9}-R^{7}$$

$$R^{8}$$

$$R^{7}$$

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in which at least one of R⁶, R⁷ and R⁸ is selected from X², and X² substituted C₁₋₄ alkyl and any others are H or C₁₋₄ alkyl; R⁹ is selected from H, C₁₋₄ alkyl, X² and X² substituted C₁₋₄-alkyl;

m is 0 or 1;

n is 1 or 2;

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X¹ is a halogen atom, a hydroxyl group, a C₁₋₆ alkoxyl group, an aryloxy group or an acyloxy group; and

 $\rm X^2$ is a halogen atom, a hydroxyl group, a $\rm C_{1-6}$ alkoxyl group, an aryloxy group or an acyloxy group;

provided that at least one of R1 to R4 is a group of formula II.

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- 2. A compound according to claim 1 in which R¹ and R² are each a group of formula II.
- 3. A compound according to claim 1 in which R^1 is a group of formula II and R^2 is $NHR^0N(R^5)_2$.
- 4. A compound according to claim 3 in which each R⁵ is the same and is H or CH₃.
 - 5. A compound according to any of claims 2 to 4 in which R¹ is at position 4 in the anthraquinone ring system and R² is in position 1.
- 6. A compound according to any preceding claim in which R³ and R⁴
 10 are selected from H and hydroxyl.
 - 7. A compound according to claim 6 in which R³ and R⁴ are both hydroxyl and are substituted at positions 5 and 8 in the anthraquinone ring system.
 - 8. A compound according to claim 6 in which R³ and R⁴ are both H.
 - 9. A compound according to any preceding claim in which m is 1.
 - 10. A compound according to any of claims 1 to 8 in which m is 0.
 - 11. A compound according to any preceding claim in which n is 2.
 - 12. A compound according to any preceding claim in which X^2 is a halogen atom or a leaving group.
 - 13. A compound according to claim 12 in which X^2 is chlorine.
 - 14. A compound according to any preceding claim in which either
 - i) R⁶ is CH₂X³ and R⁷ is H; or
 - ii) R^6 is H and R^7 is X^3

in which X^3 is a halogen atom or a leaving group.

- 15. A compound according to claim 14 in which R⁶ is CH₂X³ and R⁷ is H.
- 16. A compound according to claim 15 in which n is 2 and R^9 is CH_2X^3 in which X^3 is the same as X^3 in R^6 .
- 17. A compound according to claim 9 or claim 10 and/or claim 12 for use in a method of treatment of an animal by therapy.
 - 18. A composition comprising a compound according to claim 9 or

claim 10 and/or claim 12 and an excipient.

- 19. A composition according to claim 18 which is a pharmaceutical composition and in which the excipient is a pharmaceutically acceptable excipient.
- 20. Use of a compound according to claim 9 or 10 and/or claim 12 in the manufacture of a medicament for use in the treatment of an animal by therapy.
 - 21. Use according to claim 20 in which the animal is a human.
- 22. Use according to claim 20 or claim 21 in which the animal is suffering from a tumour and the therapy is anti-tumour therapy.
- 23. Use according to claim 22 in which the compound is a compound according to claim 9 and in which the therapy additionally involves administration of a cytotoxic agent and/or radio therapy of the tumour.
 - 24. A synthetic method in which a compound of the formula III

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in which R^{11} to R^{14} are each selected from H, X^4 , hydroxyl, C_{1-4} alkoxy, acyloxy, a group -NHR¹⁰N(R^{15})₂ in which R^{10} is C_{1-12} alkane diyl and each R^{15} is H or optionally substituted C_{1-4} alkyl, and in which X^4 is a halogen atom or a leaving group provided that at least one of R^{11} to R^{14} is X^4 ;

is reacted with a cyclic aminoalkylamine compound of the general formula $\ensuremath{\mathsf{IV}}$

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such that the group X^4 is replaced in a nucleophilic substitution reaction by a group of formula V

$$\begin{array}{c}
\begin{pmatrix} O \\ A \\ P \end{pmatrix} \\
R^{19} \\
R^{18}
\end{array}$$

$$\begin{array}{c}
R^{16} \\
R^{17}
\end{array}$$

in which either at least one of R^{18} , R^{17} and R^{18} is selected from X^5 and X^5 substituted C_{1-4} alkyl and any others are H or C_{1-4} alkyl, and R^{19} is selected from H, C_{1-4} alkyl, X^5 and X^5 substituted C_{1-4} alkyl

 X^5 is hydroxyl or a protected hydroxyl, or X^5 is a leaving group or a halogen atom different to X^4 and q is 0 or 1.

- 25. A method according to claim 24 in which at least one group X⁵ is hydroxyl or protected hydroxyl and in which the product is reacted with a halogenating compound optionally after deprotection to replace the or each X⁵ hydroxyl group by a halogen atom.
- 26. A method according to claim 25 in which the halogenating agent is a chlorinating agent.
- 27. A method according to any of claims 24 to 26 in which q is 0 and the product is oxidised at the ring nitrogen atom to form the corresponding amine oxide (q is 1).
- 28. A method according to any of claims 24 to 28 in which one of R^{11} to R^{14} is a group -NH $R^{10}N(R^{15})_2$ and which involves the preliminary step of reacting a precursor compound in which the corresponding group X^6 where X^6 is a halogen atom or a leaving group, with an acyclic aminoalkylamine compound of general formula VI

$$H_2NR^{10}N(R^{15})_2(VI)$$

in a preliminary nucleophilic substitution reaction in which X^6 is replaced by the group -NHR¹⁰N(R¹⁵)₂, in which R¹⁵ is H or an optionally substituted C₁₋₄ alkyl group.

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- 29. A method according to any of claims 23 to 26 in which R^{11} and R^{12} are the same and are X^5 and in which 2 equivalents of the cyclic aminoalkylamine compound IV are reacted whereby both groups X^4 are replaced by the said group of general formula V.
 - 30. A compound of the general formula VII

$$R^{29}$$
 R^{29}
 R^{29}
 R^{27}
 R^{28}
 R^{27}

in which R^{20} is a C_{1-12} -alkanediyl group and either R^{26} is CH_2CI , and R^{27} is H, or R^{26} is H and R^{27} is CI;

15 R^{29} is H or is the same group as R^{26} ; the or each R^{28} is H or is the same group as R^{27} ; and r is 1 or 2.

- 31. A compound according to claim 30 in which R^{20} is $(CH_2)_2$.
- 32. A compound according to claim 30 or claim 31 in which R^{26} is CH_2CI , R^{27} is H and R^{29} is selected from H and CH_2CI .
 - 33. A compound according to claim 30 or claim 31 in which R^{26} is H, R^{27} is CI, R^{29} is H and R^{28} is H.
 - 34. A compound according to any of claims 30 to 33 in which r is 1.
 - 35. A compound according to any of claims 30 to 33 in which r is 2.
 - 36. A method of synthesis of a compound as claimed in claim 30 in which a hydroxyl-substituted cyclic tertiary amine of the general formula VIII

$$\begin{array}{c|c}
 & R^{21} \\
 & R^{24} \\
 & R^{23}
\end{array}$$
(VIII)

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in which R²⁰ and r are as defined in claim 30
either R²¹ is CH₂OH and R²² is H
or R²¹ is H and R²² is OH;

R²⁴ is H or is the same group as R²¹
the or each R²³ is H or is the same group as R²²;
is amine-group protected, is then chlorinated by a process in which the OH is replaced by CI, and is deprotected to afford the desired compound of

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formula VII.